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Sep 25, 2015 · The **primary** therapy for **renal synovial sarcoma** is radical surgical resection. However, no consensus has been reached on the adjuvant treatment since at present only **case** reports have been published. The current study reports **a case** of **primary** SS (PSS) of the **kidney** and presents a review of the relevant literature.

Cited by: 4 **Author:** Zijian Wang, Zhaohui Zhong, Liang Zhu, ...
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Primary synovial sarcoma of the **kidney**: **A case report** ZIJIAN WANG, ZHAOHUI ZHONG, LIANG ZHU, WEI XIONG, CIZHONG PAN, XINJUN WANG, ZHICHAO HUANG and XIAOKUN ZHAO Department of Urology, The Second Xiangya Hospital, Central South University, Changsha, Hunan 410011, P.R. China Received September 19, 2014; Accepted April 28, 2015

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Authors: Zijian Wang · Zhaohui Zhong · Liang Zhu · Wei Xiong · Cizhong Pan · Xinjun Wang
Affiliation: Central South University
About: Oncology · Gene · Cell · **Synovial sarcoma** · Apoptosis · Molecular medicine

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May 11, 2018 · **Primary synovial sarcoma** of the **kidney** is a rare type of soft tissue **sarcoma**. Its presenting features can resemble those of other **renal** tumors; rendering its early diagnosis, a dilemma. Several cases of **renal synovial sarcoma** have been reported in the literature with varying treatment options and outcomes. This article describes a rare **case** of **primary renal synovial sarcoma** and ...

Author: Alissar El Chediak, Deborah Mukherji,... **Publish Year:** 2018

Primary synovial sarcoma of the kidney: A case report

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Sep 25, 2015 · Introduction. **Synovial sarcoma** (SS) originates primarily in the soft tissues, generally in the para-articular regions of the extremities. These tumors can, however, involve other unusual locations (), such as the pleura, lungs, mediastinum and kidneys.**Primary renal** SS is a rare carcinoma that was first described by Argani et al in 1999 ().This tumor presents a diagnostic dilemma, as it is ...

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Primary renal synovial sarcoma: A case report

Huai-Jie Cai, Nan Cao, Wei Wang, Fan-Lei Kong,Xi-Xi Sun

Abstract

Background: Synovial sarcoma is a rare mesenchymal tumor with unclear histological origin and differentiation direction, accounting for 6%-10% of soft tissue tumors. It is mainly located near the joints and tendons of the limbs, mostly occurring in children or young adults[1].Primary renal synovial sarcoma (PRSS) is very rare, accounting for approximately 1% of synovial sarcoma. It is a spindle cell tumor of mesenchymal tissue with morphological, genetic and clinical characteristics and a certain degree of epithelial differentiation. It is highly malignant, ranking the fourth highest incidence of soft tissue sarcoma[2].Here, we report the case of PRSS and share some valuable information about the disease.

Case summary: A 54-year-old male patient was admitted to the hospital due to "right kidney occupying for 2 days upon physical examination". Clinical manifestations: the patient has no cold or fever, and no frequency, urgency or pain of urination,

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Primary renal synovial sarcoma: A case report

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Renal synovial sarcoma is a recently reported neoplasm rarely seen. We **report** a **case** of **primary renal synovial sarcoma**. The signs and symptoms are similar to any **primary renal** tumour diagnosis is clinically difficult through general survey or multiple imaging modalities and requires specific molecular and genetic testing.

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